Decomposition pathways in age hardening of Ti-Al-N films

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The ability to increase the thermal stability of protective coatings under work load gives rise to scientific and industrial interest in *age hardening* of complex nitride coating systems such as ceramic-like $\text{Ti}_{1-x}\text{Al}_x\text{N}$. However, the decomposition pathway of these systems from single-phase cubic to the thermodynamically stable binary nitrides (cubic TiN and wurtzite AlN), which are essential for *age hardening*, are not yet fully understood. In particular, the role of decomposition kinetics still requires more detailed investigation. In the present work, the combined effect of annealing time and temperature upon the nano-structural development of $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$ thin films is studied, with a thermal exposure of either 1 min or 120 min in 100 °C steps from 500 °C to 1400 °C. The impact of chemical changes at the atomic scale on the development of micro-strain and mechanical properties is studied by post-annealing investigations using X-ray diffraction, nanoindentation, 3D-atom probe tomography and high-resolution transmission electron microscopy. The results clearly demonstrate that the spinodal decomposition process, triggering the increase of micro-strain and hardness, although taking place throughout the entire volume, is enhanced at high diffusivity paths such as grain or column boundaries and followed within the grains. *Ab initio* calculations further show that the early stages of wurtzite AlN precipitation are connected with increased strain formation, which is in excellent agreement with experimental observations. © 2011 American Institute of Physics. [doi:10.1063/1.3610451]

I. INTRODUCTION

The combination of high hardness, good oxidation resistance and the possibility to tune the Ti/Al ratio and micro-structure of $\text{Ti}_{1-x}\text{Al}_x\text{N}$ thin films in a wide range, e.g., by different deposition processes, makes $\text{Ti}_{1-x}\text{Al}_x\text{N}$ one of the most versatile protective hard coating materials for many applications and substrates.¹ The major use as protection for drills, tools, dies, and molds benefits from the high thermal stability of $\text{Ti}_{1-x}\text{Al}_x\text{N}$, which is known to be strongly correlated with its microstructure and chemical composition.² Due to typical substrate temperatures of ~500 °C (~0.2–0.3 of the melting temperature), the atomic reassembly during condensation from the vapor phase is limited, which results in a meta-stable solid solution of $\text{Ti}_{1-x}\text{Al}_x\text{N}$. During common physical vapor deposition (PVD) processes, crystallization in the cubic (c) NaCl (B1) structure is usually found up to Al contents of the metallic sublattice $x \sim 0.6$ –0.7.^{3,4} Higher Al contents favor the wurtzite (w) ZnS (B4) structure. The metastable phases obtained from these processes exhibit

a driving force for separation toward their stable constituents c-TiN and w-AlN, strongly scaling with the Al content.^{5–9}

In common milling operations, the high thermal work load on the coated tools typically reaches temperatures in excess of ~900 °C. This enables sufficient diffusion in the coating to induce the chemically driven isostructural decomposition of metastable $\text{Ti}_{1-x}\text{Al}_x\text{N}$ into TiN- and AlN-enriched domains.¹⁰ In many reports^{2,8,11–13} this effect is attributed to spinodal decomposition of the c- $\text{Ti}_{1-x}\text{Al}_x\text{N}$ matrix, resulting in *age hardening* of $\text{Ti}_{1-x}\text{Al}_x\text{N}$.^{8,14}

In addition to the chemical driving force (due to a positive mixing enthalpy over the whole composition range of $\text{Ti}_{1-x}\text{Al}_x\text{N}$), also the development of elastic strain energy and the formation of new interfaces have to be taken into account.¹² Outside the spinodal range, $0.27 < x < 0.98$,^{9,15–17} at 900 °C, precipitation by nucleation and growth is the dominating reaction for decomposition into c-TiN and w-AlN, while within the spinodal range the iso-structural decomposition (without nucleation barrier) to form TiN- and AlN-enriched domains, is the favored pathway.

Although the influence of annealing temperature on the structural evolution of $\text{Ti}_{1-x}\text{Al}_x\text{N}$ thin films was investigated by computational^{12,15,16,18,19} and experimental^{8,11,12,14} methods, a combinatorial description of the kinetics and decomposition

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pathway is of paramount importance. In the present study, a detailed analysis of the structural and chemical evolution of c-Ti_{1-x}Al_xN thin films as a function of temperature and time is presented and related to the development of the mechanical and electrical properties. The assessment of coating structure by X-ray diffraction (XRD) and high resolution transmission electron microscopy (HR-TEM) is combined with a chemical and morphological analysis using three-dimensional atom probe tomography (3D-APT) and correlated to nanoindentation and electrical resistivity experiments. *Ab initio* calculations were performed to facilitate the interpretation of the experimental findings. The present study aims to elucidate the phase development, especially the morphology of spinodal decomposition, of metastable single phase c-Ti_{1-x}Al_xN toward its stable constituents c-TiN and w-AlN.

II. EXPERIMENTAL

A. General considerations

Two different polycrystalline substrates, low alloy steel foil (thickness 0.2 mm) and polycrystalline Al₂O₃ platelets (20 × 7 × 0.5 mm) were selected for the present investigations. The low alloy steel foil can easily be removed chemically to avoid substrate interference during time and temperature dependent 3D-APT, XRD and TEM studies, whereas the polycrystalline Al₂O₃ provides an ideal substrate for nanoindentation and resistivity measurements of the thin films. For the deposition of the Ti_{1-x}Al_xN thin films a mixed Ar/N₂-plasma discharge and a Ti_{0.50}Al_{0.50} target (Plansee AG, Ø 150 × 5 mm) were used in a lab-scale magnetron sputter deposition plant. For a more detailed description of the parameters used, see Ref. 20.

B. Experimental details

After deposition, both the free standing thin film powder (after chemical removal from the low alloy steel substrate) and the coating on Al₂O₃ were annealed in a vacuum furnace (HTM Reetz GmbH, base pressure < 5 × 10⁻⁴ Pa) at different temperatures (*T_a*) from 500 to 1400 °C in 100 °C steps, with a heating rate of 20 K min⁻¹ and a cooling rate of 50 K min⁻¹, and subsequently measured at room temperature. The reason for this was to minimize structural changes during the cooling process from *T_a*. The samples were kept at *T_a* for either 1 min or 120 min. Subsequent to the annealing treatment, the electrical resistivity of the coatings on Al₂O₃ was determined by a Jandel four-point probe. In this measurement, a current of 900 μA was used and the sample geometry corrected according to Refs. 21 and 22. For nanoindentation a UMIS nanoindenter, equipped with a Berkovich tip, was used. A plateau test²³ was performed, with stepwise increasing load from 6 to 25 mN for at minimum 20 indentations, which ensured that the indentation was contained within the upper 10% of the coating in order to minimize possible substrate interference. Structural analysis was made using X-ray diffraction with a Bruker D8 and Cu-K_α emission, applying Bragg-Brentano geometry for the powdered free standing film material and grazing incidence at 2° incidence angle for coatings on Al₂O₃ substrates. Quantitative XRD data obtained from the powdered film material were first

fitted using the PeakFIT[®] program, applying a Voigt-function for all distinguishable cubic and wurtzite peaks in the measured range between 30 to 85°, prior to a Williamson-Hall analysis.²⁴ In preparation for TEM and EDX analysis (using a C_s-corrected JEOL 2100 F equipped with a Gatan energy dispersive X-ray spectrometer (EDX) and operated at 200 kV), plan-view samples were polished and ion-milled using Argon ions in a Gatan Precision Ion Polishing System (PIPS). EDX measurements from scanning electron microscopy (SEM) studies, using a Zeiss Evo 50, of the coatings on alumina as well as the TEM investigations indicate consistent values of ~22.5 ± 0.5 at.% for Ti, ~26.5 ± 0.5 at.% for Al and ~51 ± 1 at.% for N. The coating composition normalized to 50 at.% N will therefore be referred as Ti_{0.46}Al_{0.54}N.

Atom probe tomography (3D-APT) enables sub-nanometer resolution in the measurement direction,²⁵ and provides a quantitative morphological description of the material in three dimensions. Previous work^{10,20,26} introduces this technique applying an Imago LEAP 3000X-HR to the field of hard coatings and discusses APT sample preparation, experimental parameters and limitations of the technique in more detail. The parameters utilized for the *ab initio* calculations can be found in Ref. 27.

III. RESULTS AND DISCUSSION

A. Structure and mechanical properties

Comparative X-ray analysis of powdered coating samples and coatings on Al₂O₃ after annealing for 1 min at *T_a* is shown in Figs. 1(a) and 1(b), respectively. For simplicity only the XRD patterns after annealing at 900, 1000, 1100, and 1300 °C are presented. The observed structural features, such as peak intensities and the formation of w-AlN, are in good agreement for both the powdered coating and the coating on substrate. A single phase cubic solid solution of Ti_{0.46}Al_{0.54}N, with cubic lattice parameter *a_c* = 4.181 Å, is obtained from the powdered samples in the as deposited state. The samples maintain their single phase cubic structure with increasing *T_a* up to ~1000 °C. For *T_a* between 900 and 1000 °C an asymmetric peak broadening is observed, which can be interpreted as a sign of isostructural formation of

FIG. 1. (Color online) XRD pattern of annealed (a) Ti_{0.46}Al_{0.54}N film powder and (b) Ti_{0.46}Al_{0.54}N film on Al₂O₃ substrate.

c-AlN- and c-TiN-rich domains.¹⁴ The variation of the cubic (111) and (002) peak intensities, visible predominantly in the X-ray patterns of the films on Al_2O_3 substrate, is related to the strong elastic anisotropy in $\text{Ti}_{1-x}\text{Al}_x\text{N}$ solid solutions, which scales with the Al-content on the metallic sub-lattice.²⁸ $\text{Ti}_{1-x}\text{Al}_x\text{N}$ with low Al-contents is known to be stiffer in the (002) direction, whereas high Al contents exhibit a higher stiffness in (111) direction. Already for $x \geq 0.28$ the (002) direction exhibits a lower elastic modulus than the (111) direction.²⁸ Therefore, to minimize strain energy (due to coherency between the cubic AlN-enriched domains, the remaining matrix and TiN-enriched areas) during spinodal decomposition, the orientation of the $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$ coating will change toward (002). Assuming a higher strain level in the coating on Al_2O_3 , stemming from the additional strain interaction with the substrate, results in pronounced peak intensities of the (002) reflexes, especially visible at $T_a = 1000$ and 1100°C , compared to the powdered film. This result correlates with substrate-induced compressive stresses in the films favoring the cubic phase in $\text{Ti}_{1-x}\text{Al}_x\text{N}$.²⁹

The structural evolution of c- $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$ has a remarkable impact on the mechanical properties, presented in terms of the dependence of hardness (H) and indentation modulus (E) on T_a and t_a in Fig. 2. In good agreement with earlier studies,^{8,14} a hardness increase of $\sim 25\%$, from 26 ± 2 GPa in the as deposited state to 32.5 ± 1.2 GPa after annealing for 1 min at 1000°C is observed. Similarly the hardness increases when annealing at T_a for $t_a = 120$ min, although the peak hardness of 33.3 ± 1.8 GPa (an increase of $\sim 28\%$) is obtained at a lower temperature of 800°C . For both annealing times, the hardness significantly decreases for T_a above the peak value, which is called *overaging*,³⁰ whereby the hardness decline is slower for coatings annealed for $t_a = 120$ min. As previously reported^{8,14} the hardness decrease is connected with the formation of w-AlN; see for comparison Figs. 1 and 2. The behavior of the indentation moduli is similar, but especially for $t_a = 1$ min E first slightly decreases with T_a before the decomposition process initiates an

increase of E , which then decreases again due to the formation of w-AlN in the *overaging* regime.

In order to effectively eliminate substrate influence, for a further quantitative study by means of Williamson-Hall analysis,²⁴ only the powder XRD-data were evaluated for all annealing treatments. To give an overview of the phase development between the samples annealed for 1 and 120 min, *quasi* two-dimensional plots of the recorded XRD-patterns are given in Figs. 3(a) and 3(b), respectively, where the X-ray intensities are illustrated by a corresponding color code. The XRD patterns were recorded in 100°C steps after annealing and are linearly interpolated. Significant peak broadening indicates phase separation of the single phase cubic matrix at temperatures below 1100°C for $t_a = 1$ min [Fig. 3(a)] and 900°C for $t_a = 120$ min [Fig. 3(b)]. The peak broadening of the cubic reflexes, as discussed above and in literature,^{8,14} results from spinodal decomposition, where TiN- and AlN-rich domains are formed that have slightly different lattice parameters. Due to the diffuse nature of the domain boundaries¹⁰ no sharp signal of the two domains can be detected. For both annealing times it was found that w-AlN can be detected first with its (10–10) peak at $\sim 33.2^\circ$ at $T_a = 1100^\circ\text{C}$ for $t_a = 1$ min and $T_a = 900^\circ\text{C}$ for $t_a = 120$ min, followed by higher indexed reflexes at higher T_a . As a guide for interpretation, a solid orange line connects the signals at 2% of the highest detected intensity, highlighting the discussed XRD-peak broadening and the advent of the (10–10) reflex for w-AlN. By increasing the annealing time from 1 to 120 min, the formation of w-AlN is shifted to $\sim 200^\circ\text{C}$ lower temperatures. This indicates a strong diffusion-driven and hence time-dependent decomposition behavior at different temperatures, confirming earlier predictions.¹²

The development of the lattice parameters as a function of T_a and t_a is highlighted in Fig. 4. The cubic lattice parameter a_c was calculated from the main XRD response at $\sim 43^\circ$, (002) peak in c- $\text{Ti}_{1-x}\text{Al}_x\text{N}$. With increasing T_a the cubic lattice parameter decreases for the 1 min and 120 min annealing to a minimum of ~ 4.16 Å at $T_a = 1000^\circ\text{C}$ and 800°C , respectively, coinciding with the maximum in H and E , and then increases to meet the value of 4.24 Å for TiN; see Fig. 4. Similarly, the hexagonal lattice parameters a_w and c_w were determined from w-AlN (10–10) and (0002) peaks, respectively. At high T_a , where w-AlN can be detected, both a_w and c_w deviate significantly from the stoichiometric values of 3.11 and 4.98 Å for w-AlN³¹ (see Fig. 4).

The impact of this structural development on the microstrains (ϵ) of the cubic phase is given in Fig. 5(a). Please note that these strains refer only to intrinsic stresses of second and third order, as a result of the incorporation of defects³² and initial chemical fluctuations of Ti- and Al-concentrations¹⁰ because the measurements are conducted on free-standing film material (see Sec. II B). Therefore, only stresses between individual grains or columns (second order) and within these microstructural features (third order) are responsible for the observed microstrains. The initial strain of the c- $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$ solid solution remains at $\sim 0.8\%$ with annealing up to the deposition temperature of 500°C . The samples annealed for 1 min at T_a exhibit a moderate increase of ϵ with T_a , which suggests an increase of coherency strain as only isostructural decomposition is expected in this

FIG. 2. (Color online) Hardness (H) and indentation modulus (E) of $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$ as a function of T_a for $t_a = 1$ min (full symbols) and $t_a = 120$ min (open symbols).

FIG. 3. (Color online) Quasi-two dimensional XRD-plot of $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$ powder annealed to T_a for $t_a = 1$ min (a) and $t_a = 120$ min (b). Note that the logarithmic intensity corresponds to the color code in the scale bar. The orange line highlights the 2% threshold value of the normalized intensity.

temperature range; compare to Figs. 1 and 3. The atomic assembly due to isostructural decomposition is schematically presented in Fig. 5(b). For the samples annealed for 120 min at T_a there is a more pronounced increase of ε with T_a , indicating the strong impact of t_a on the decomposition kinetics. The strain values after annealing for $t_a = 1$ and 120 min at $T_a = 1000$ and 800°C , are 1.31 and 1.33%, respectively. This

means a strain increase of $\sim 60\%$ compared to the as deposited value, where the hardness is at its maximum of 32.5 ± 1.2 GPa ($t_a = 1$ min) and 33.3 ± 1.8 GPa ($t_a = 120$ min), compare to Fig. 2, and the cubic lattice parameter is at its minimum of ~ 4.16 Å, see Fig. 4.

The decrease of a_c is thus not simply related to the recovery of built-in defects, but rather an effect of isostructural decomposition and the concomitant development of microstrains, as schematically shown in Fig. 5(b) for the single phase regime. The maximum microstrain of 1.75 and 2.00% for $t_a = 1$ and 120 min is observed at $T_a = 1100$ and 900°C , respectively, and consequently 100°C above the temperature for peak hardness and minimum cubic lattice parameter. Here, already small fractions of w-AlN with distorted lattice parameters a_w and c_w can be detected, as evident in a comparison between Fig. 3 and Fig. 4. Within the dual phase regime, ε declines more strongly in the samples annealed for $t_a = 1$ min compared to those annealed for $t_a = 120$ min which exhibit a smaller strain decrease between 900 and 1100°C before ε strongly decreases at higher T_a . This strain development is in excellent agreement with the observed hardness values, which decrease more slowly for the samples annealed for $t_a = 120$ min in the temperature range ($900^\circ\text{C} < T_a < 1100^\circ\text{C}$); compare Fig. 2. Thus almost similar hardness values were obtained for $t_a = 1$ and 120 min at $T_a = 900$ and 1100°C , although stemming from two significantly different decomposition states with respect to strain and structure (e.g., before and after precipitation of w-AlN). A further increase of T_a (as soon as w-AlN precipitates) results in coarsening of the dual phase structure and a concomitant strain decrease for both t_a , as shown schematically in Fig. 5(c) and described in the following section.

FIG. 4. (Color online) Lattice parameters of the cubic (a_c - above) and the wurtzite phase (a_w , c_w - below) as obtained from powder X-ray diffraction. Full and open symbols refer to the values obtained after annealing for $t_a = 1$ min and 120 min, respectively. Note that the error bars are only visible when they are not smaller than the symbol size.

B. Three-dimensional-morphology and chemical assembly

For additional chemical information at the atomic scale during the decomposition process, atom probe measurements

FIG. 5. (Color online) (a) Microstrain, ϵ , obtained from the powder samples after annealing at T_a for $t_a = 1$ min (black full symbols) and $t_a = 120$ min (gray open symbols) using the Williamson-Hall-method.²⁴ Schematic illustration of the atomic assembly for (b) the isostructural decomposition into c-TiN- and c-AlN-enriched domains which causes the strain increase in the single phase regime, and (c) the dual phase structure at higher T_a resulting in strain decrease due to grain growth.

were performed on samples annealed at 900 °C for $t_a = 1$ and 120 min. These conditions were chosen, as they represent samples that are 100 °C below (for $t_a = 1$ min) and 100 °C above (for $t_a = 120$ min) those with observed peak hardness. Additionally, a sample in the as deposited state (for reference) and a sample annealed at $T_a = 1350$ °C for 1 min are investigated. The latter represents the decomposed state of the film composed of c-TiN and w-AlN as shown in Figs. 1 and 3. Statistical 3D-APT data treatment in combination with 2 D-concentration plots and isoconcentration surfaces were selected to graphically portray the morphological evolution of the c-Ti_{0.46}Al_{0.54}N films under thermal load.

The datasets were evaluated according to a maximum likelihood method, using the Langer-Bar on-Miller (LBM) treatment,^{33,34} which approximates the probability distributions of the decomposing phases as two Gaussian distributions of equal width, σ , located at respective concentrations μ_1 and μ_2 . Thereby, the magnitude of the average local chemical fluctuations of the metallic sublattice can be expressed as the difference between the two Gaussians, $\mu_2 - \mu_1$, as shown in Fig. 6(a). Since the magnitude of compositions in APT-data is extremely sensitive to the selected

block size,³⁵ the LBM-treatment was carried out for different block sizes as presented in Fig. 6(a). Too small block sizes lead to statistical noise in the calculation and tend to overestimate the magnitude of $\mu_2 - \mu_1$, reflected by a decreasing slope with increasing block size up to ~ 100 atoms/block. This regime is followed by block sizes of the same order as the chemical composition before too large block sizes smooth out the real chemical fluctuations³³ and $\mu_2 - \mu_1$ rapidly increases again. It can be seen that the difference in chemical composition ($\mu_2 - \mu_1$) of the samples in the as deposited and annealed state reaches a first plateau at a block size of ~ 100 atoms/block. Consequently, this regime was chosen for the determination of the LBM statistics, which are presented in Table I.

A comparison of the observed frequency distribution with a binomial distribution and the LBM-model is presented for the Al-ions in Fig. 6(b). The presentation of the obtained values for Al is favorable to Ti due to the slightly higher Al-content in the film, although the results for Ti are qualitatively similar. For all samples a significant deviation from a random solid solution, usually described as a binomial distribution, is visible and the LBM-method gives the best

FIG. 6. (Color online) (a) The difference of the two Gaussians $\mu_2 - \mu_1$, reflecting the chemical compositions of the respective TiN- or AlN-enriched domains, as obtained from the Langer-Bar on-Miller treatment of the 3D-APT data as a function of block size. (b) Frequency distributions of the Al concentration for the as deposited (top) and annealed to $T_a = 900^\circ\text{C}$ for $t_a = 1$ min (middle) and 120 min (bottom) state. The inserts present a magnification of the Al-rich part of the distribution. The green dashed line indicates the average Al composition of 26.5 at. %.

approximation for the as deposited state. In addition to an increasing width of the distribution, indicating phase separation, the samples annealed for $t_a = 1$ and 120 min at $T_a = 900^\circ\text{C}$ exhibit an increasing asymmetry, which is more pronounced for the 120 min sample. The high χ^2 value for $T_a = 900^\circ\text{C}$ and $t_a = 120$ min (Table I) suggests that the LBM-model becomes insufficient to describe the present decomposition state. These results are consistent with the

TABLE I. Parameters of the LBM-fits to the frequency distributions shown in Fig. 6.

		as deposited	900 °C – 1 min	900 °C – 120 min
μ_1	[%]	25.9	25.3	29.0
μ_2	[%]	28.9	32.3	41.0
σ	[%]	2.0	4.0	4.0
χ^2	[-]	45.9	90.1	1074
d.o.f.	[-]	35	41	50

experimentally observed asymmetric peak development of the cubic reflexes in the X-ray diffraction investigations. The small inserts in each frequency distribution highlight the deviation from the binomial (red solid line) and LBM-model (gray dashed line with circles) at high Al-concentrations. The results suggest the presence of stoichiometric AlN (Al-content of 50 at. %) for the sample annealed at 900°C for 120 min.

The graphical presentation of the investigated samples, shown in Figs. 7(a)–7(d), was performed by isosurfaces^{36,37} at Al-concentrations of 28.5 at. % (as deposited), 29.5 at. % (900°C for 1 and 120 min), and 46.5 at. % (1350°C for 1 min), respectively. These isoconcentration values are 2, 3, and 20 at. % above the 26.5 at. % average Al-concentration, in order to graphically emphasize the increasing local elemental fluctuations and phase separation with T_a and t_a . The two-dimensional concentration plots represent the respective Al and Ti concentrations, projected toward the y-direction of the highlighted yellow boxes ($50 \times 2 \times 80$ nm) in the center of the tips. Due to the sample preparation technique, the field evaporation (measurement direction z) of the shown tips was perpendicular to the thin film growth direction (y -axis) as described in Ref. 20. Hence, the 2 D-plots give plan-view information about the local element distribution across column boundaries.

Already in the as deposited state, local elemental fluctuations of Al and Ti ions in opposite directions were detected [Fig. 7(a)], confirming the deviation from an ideal random element distribution as mentioned above and presented in Ref. 10. Note the different scales for the Al and Ti concentrations in the different annealing states enhancing the color contrast of the respective decomposing domains. After annealing at 1350°C for 1 min, the formation of almost stoichiometric TiN and AlN can be confirmed by 3D-APT [see Fig. 7(d)]. Only small fractions of Al or Ti ions can be detected within the coarsened grains of the opposite species.

Annealing at 900°C for 1 min [see Fig. 7(b)] leads to an increasing compositional amplitude and the formation of TiN- and AlN-enriched domains, spanning an interconnected 3D-network compared to the as deposited state reported earlier.¹⁰ Although an increasing intensity of the compositional oscillations throughout the whole volume can be found, a stronger Al-enrichment can be detected at grain boundaries, highlighted by the red arrow. The development of a 3D-interconnected structure of c-Ti_{1-x-Δ}Al_{x+Δ}N and c-Ti_{1-x+Δ}Al_{x-Δ}N is a result of the spinodal decomposition process and therefore coincides with the structural investigations as described

FIG. 7. (Color online) Isoconcentration surfaces for Al contents drawn at the given threshold values for the as deposited state (a), after annealing at 900 °C for 1 min (b) and 120 min (c) and after annealing at 1350 °C for 1 min (d). The 2 D-concentration plots on the right side present the respective element distribution of Al or Ti calculated in the y-direction (initial growth direction) of the box shown in the center of the tips. The arrows indicate regions with highest Al concentration gradients identified as column boundaries, see also Fig. 9.

above. Using electron energy loss spectroscopy (EELS) Knutsson *et al.* observed a qualitatively similar element distribution in arc-evaporated TiN-Ti_{0.34}Al_{0.66}N multilayer coatings after annealing to 900 °C.^{38,39} The locally higher compositional amplitude at former grain boundaries is likely to originate from the higher diffusivity as compared to the grain interior. Also a Kirkendall-like behavior, due to the higher diffusivity of Al compared to Ti, can contribute to this effect as proposed by Johnson *et al.* for Ti_{0.34}Al_{0.66}N after

annealing at 900 °C.²⁶ The sample annealed at 900 °C for 120 min [Fig. 7(c)] indicates that the compositional mismatch of the TiN- and AlN-enriched domains within the grains significantly increased compared to samples annealed only for 1 min. Also here the highest Al contents were found at the grain boundaries (red arrow). At this annealing stage, the AlN-rich domains are clearly elongated along the growth direction of the film and locally reach an Al concentration of ~50 at.%; compare also Fig. 6(b). Theoretical and experimental

FIG. 8. (Color online) *Ab initio* results in the change of the total internal energy of w-AlN due to Ti-impurities ($\Delta U_{\text{comp.}}$ – red full symbols) as a function of Al-mole fraction (x) and the corresponding increase of the atomic volume. The total internal energy changes due to strain (ΔU_{strain} – open black symbols) of stoichiometric w-AlN distorted to the same volumes reaches a significantly lower increase in internal energy and is thus suggested to be energetically favorable.

investigations on $\text{Ti}_{0.50}\text{Al}_{0.50}\text{N}$ thin films^{40,41} suggest a different growth behavior of the TiN- or AlN-rich domains in the growth direction and perpendicular to it due to the strong dependence of the mixing enthalpy on the local atomic ordering and strain effects.⁴² A further argument takes into account the generation and reduction of internal stresses during decomposition which stems from a strong orientation relationship between the decomposing counterparts.⁴³

In order to determine the impact of strain or Ti-impurities on the total internal energy of w-AlN, *ab initio* calculations were performed. Figure 8 shows that the change in internal energy, $\Delta U_{\text{comp.}}$, of w-AlN strongly increases to 0.17 eV at^{-1} if Ti atoms substitute Al in w-AlN (red full squares) to obtain a 4.3% larger cell size. If, on the other hand, stoichiometric w-AlN is strained by 1.43% to reach a comparable cell size, the change in internal energy, ΔU_{strain} , (black open squares) is much smaller ($\sim 0.015 \text{ eV at}^{-1}$), see Fig. 8. This result is in general agreement with the limited solubility of Ti in w-AlN and suggests that the lattice parameter deviation of w-AlN during precipitation, as obtained from our experiments (Figs. 4 and 5), is mainly caused by strain rather than chemical variation.

More detailed investigations of the atomic arrangement at the grain boundaries are made using TEM. Figure 9(a) shows a bright-field (BF) plan view image of the coating annealed at 900°C for 120 min. The insert shows the corresponding selected area electron diffraction (SAED) pattern, indicating strong c-Ti_{1-x}Al_xN reflexes and only little intensity of the w-AlN phase, in line with the X-ray investigations. The corresponding dark-field image, highlighting regions obtained from the w-AlN (10–10) reflex indicated in the SAED pattern, is shown in Fig. 9(b). Regions corresponding to the w-AlN phase are especially visible at the grain boundaries. The HR-TEM image, Fig. 9(c), from the area indicated with a rectangle in the dark-field image [Fig. 9(b)] emphasizes the presence of

FIG. 9. (Color online) (a) Plan-view BF-TEM micrograph of the $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$ thin film annealed to 900°C for 120 min, the insert shows a SAED pattern of this region. (b) Dark field TEM image from the wurtzite reflex indicated in the SAED pattern. (c) HR-TEM image of the highlighted region from (b) showing w-AlN at the grain boundaries. The FFT inserts were obtained from the indicated regions 1 and 2, respectively. (d) The elastic anisotropy of w-AlN shown for the change of E -modulus with respect to the $[0002]_{\text{w-AlN}}$ direction. Note that $[0002]$ and $[10-10]$ directions refer to a deformation along the c - and a -axis of the w-AlN, respectively.

semicoherent and incoherent interfaces between the w-AlN precipitates and the surrounding cubic $\text{Ti}_{1-x}\text{Al}_x\text{N}$ matrix. The measured lattice constant a_c of the c- $\text{Ti}_{1-x}\text{Al}_x\text{N}$ matrix varies between 4.18 Å and 4.21 Å, depending on the area of investigation. This modulation corresponds to the local Al-fluctuations in agreement with literature.⁴⁴ Fast Fourier transformation (FFT) of region 1, corresponding to the w-AlN precipitate, exhibits three sets of additional reflexes compared to the solely cubic reflexes of region 2 [see inserts in Fig. 9(c)]. Since the w-AlN precipitate is highly strained, the reflexes overlap with the cubic reflexes in one direction, indicating a partial coherency of the $(01-1-1)_{\text{w-AlN}}// (002)_{\text{cubic}}$ planes, whereas the other two are in no orientation relationship with the surrounding matrix. Rafaja *et al.*⁴³ recently proposed an orientation relationship between $(11-1)_{\text{cubic}}// (0002)_{\text{w-AlN}}$ planes in as deposited nanocomposite $\text{Ti}_{0.50}\text{Al}_{0.50}\text{N}$ coatings, which exhibits the lowest mismatch between the two phases along the $[1-10]_{\text{cubic}}// [10-10]_{\text{w-AlN}}$ directions. However, the existence of the partial coherency also along other crystallographic orientations can be explained by the high distortion of the w-AlN in its initial precipitation stage, which has to compensate the $\sim 24\%$ higher volume per formula unit of w-AlN compared to c-AlN.⁴⁵ Further, the elastic anisotropy of w-AlN, regarded as a function of the angle with respect to the $[0002]_{\text{w-AlN}}$ direction, is shown in Fig. 9(d). The lower elastic modulus in a direction perpendicular to the $(01-1-1)_{\text{w-AlN}}$ plane therefore suggests the easier displacement of $(01-1-1)$ planes, compared to the $(10-10)_{\text{w-AlN}}$ planes, in agreement with the FFT of region 1. The observed strain maximum in the cubic phase (see Fig. 5) therefore results from the w-AlN precipitation at this annealing state.

C. Electrical resistivity

The sheet resistivity of $\text{Ti}_{1-x}\text{Al}_x\text{N}$ coatings is known to strongly scale with the Al mole fraction, x , and the structure obtained after deposition.⁴⁶ TiN films exhibit an almost metallic

FIG. 10. Electrical resistivity of the $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$ thin films measured after annealing at T_a and $t_a = 1$ min (black full symbols) and 120 min (gray open symbols). The evolution of sheet resistivity reflects the state of decomposition (compare Figs. 3 and 7).

conduction behavior ($\sim 0.1 \text{ m}\Omega \text{ cm}$ ⁴⁷), while AlN films are semiconducting ($\sim 10^{13}$ to $10^{15} \text{ m}\Omega \text{ cm}$ ⁴⁸). Due to the thermally induced structural changes of c- $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$, the sheet resistivity decreases by 1 order of magnitude from $\sim 4.6 \text{ m}\Omega \text{ cm}$ after deposition to $\sim 0.5 \text{ m}\Omega \text{ cm}$ after annealing at $T_a = 1400^\circ\text{C}$ for 1 min and $\sim 0.2 \text{ m}\Omega \text{ cm}$ for 120 min, respectively; see Fig. 10. As suggested above, the spinodal morphology of c- $\text{Ti}_{1-x}\text{Al}_x\text{N}$ and c- $\text{Ti}_{1-x}\text{Al}_x\text{N}$ domains, spanning an interconnected 3D-network throughout the volume of the film, opens a percolation pathway for charge transfer within the TiN-enriched domains (Fig. 7), being the electrically better conductive part of the composite. The state of decomposition is therefore reflected by the sheet resistivity, which goes along with the structural investigations as a function of T_a and t_a .

IV. CONCLUSIONS

The present work deals with the pathway of decomposition from single phase c- $\text{Ti}_{1-x}\text{Al}_x\text{N}$ toward a dual phase structure composed of c-TiN and w-AlN, involving spinodal decomposition and subsequent precipitation and growth. We employ the model system $\text{Ti}_{1-x}\text{Al}_x\text{N}$ with an Al content of 54 at.% at the metallic sublattice as this provides a huge decomposition driving force deep in the spinodal region, combined with a single phase cubic structure of large columnar grains after deposition. Three-dimensional atom probe tomographic studies of free standing coating samples annealed at 900°C show that the isostructural decomposition of c- $\text{Ti}_{0.46}\text{Al}_{0.54}\text{N}$ occurs within the entire coating volume. However, a higher AlN-enrichment was found at preferred diffusion pathways such as column or grain boundaries, while the extent of the local TiN- and AlN-enrichment in the grain interior is slightly retarded and becomes more pronounced with increasing annealing time from 1 to 120 min. At this stage, first highly distorted w-AlN domains at the grain boundaries can be detected, as shown by HR-TEM investigations. Concomitant X-ray and nanoindentation experiments show that the hardness maximum of $32.5 \pm 1.2 \text{ GPa}$ ($t_a = 1 \text{ min}$, $T_a = 1000^\circ\text{C}$) and $33.3 \pm 1.8 \text{ GPa}$ ($t_a = 120 \text{ min}$, $T_a = 800^\circ\text{C}$) is achieved in the single phase regime where the relative microstrain shows an increase of $\sim 60\%$. The results further demonstrate that the peak hardness is obtained at $\sim 100^\circ\text{C}$ lower temperatures than the peak strain, which is observed during the early stages of w-AlN precipitation. The observed precipitation of highly strained w-AlN is supported by *ab initio* calculations, exhibiting a significantly higher change in internal energy due to residual Ti-impurities ($\Delta U \sim 0.17 \text{ eV at}^{-1}$) than to strain ($\Delta U \sim 0.015 \text{ eV at}^{-1}$) for a comparable volume change of the cell by $\sim 4\%$. The subsequent growth of w-AlN grains and out-diffusion of Al from the c-TiN-enriched domains with further increase of T_a , results in a loss of coherency and causes a significant strain and hardness decrease. Moreover, the presented structural evolution goes in line with a reduction in electrical resistivity by 1 order of magnitude, which is related to the formation of a 3D-interconnected network of c-TiN-enriched domains (and finally c-TiN) with essentially lower electrical resistivity.

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